NEW YORK UNIV N Y DEPT OF PHYSICS
PULSE AREA EFFECTS IN RESONANT MULTIPHOTON IONIZATION, (U)
SEP 79 E J ROBINSON
N00014-77-C-0553 F/6 7/4 AD-A075 831 UNCLASSIFIED . NL END DATE FILMED | OF | AD A075831

Submitted to J. Phys. B





AD A 0 75831

Pulse Area Effects in Resonant Multiphoton Ionization

E.J./Robinson

Department of Physics

New York University

4 Washington Place

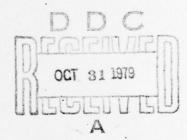
New York, N.Y. 10003 USA

11 Sep 79

(12) 8 /

DISTRIBUTION STATEMENT And Approved for public releases
Distribution Unlimited

Sept 1979



Reproduction in whole or in part is permitted for any purpose of the United States Government.

Supported by the U.S. Office of Naval Research under Contract No. 100014-77-C-0553.

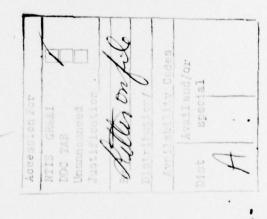
406 850

79 09 17 077

DOC FILE COPY

bstract

A theory of resonant multiphoton ionization by emooth pulses is developed. It is shown that the spectrum of the effect may exhibit structure associated with time domain correlations.



over intermediate states. In the crudest approximation, the theory for these processes typically requires the evaluation of summetions on adiabatically. If the intensity is low and there are no nearly lowest order of time-dependent perturbation theory consistent with Among the effects of nomineer optics, multiphoton ionization assumes that the radiation field is monochromatic and is switched resonant intermediate states, the transition rate is given by the binding energy. Much work has been done on this problem, and it remains an active area of research (Plank et al 1976, Flank and Copfenstein 1965, Eebb 1966, 1967, Bebb and Gold 1966, Chin and the invention of the laser (Celtman 1963, 1965, Eall et al 1965, the number of photons of energy hy needed to compensate for the and detachment were among the earliest to be investigated after Isenor 1967, Chin et al 1968, Young et al 1969, Fox et al 1971, Kogan et al 1971, Agostíni et al 1971, Inger and Robinson 1970, absorption of more than one quantum. The calculation of rates Eall 1966, Robinson and Geltman 1967, Zernik 1964, Zernik and These are bound-free electronic transitions in atomic systems generated by the simultaneous Rachman 1975, Teague et al 1976.) Lurein and Mainfray 1972).

In recent times, with the evallability of tumable laters, the study of multiphoton ionization where one or more intermediate states are close to resonance has come to the fore (Seers and Armatrong 1975, Geltman 1979, Dixit and Lembropoulous 1979, Gentler and frahin 1979, Energy and C'Scil

that these systms are excluded.) The simple perturbation theory, acceptable in the nonresonant case, becomes flowalfd when the population of the initial level is depleted and/or its phase is significantly affected by coupling to the intermediate states. Theories which take this into account have been developed and predict interesting effects such as structure in the photolonization specific interesting effects the departure of the transition probability from a simple linear function of time and bundwidth effects. The richness of this field has stimulated great activity.

coupled bound levels, but has a very different origin. It involves the AC Stark effect in experiments where there are three resonantly twice the area used in the sense of photon echoes.) In principle, resonance (ISB), exhibits maxima as the laser is detuned above and below, and then falls off rapidly. This superficially resembles the radiation from only one pulsed laser, not two. I: addition, going population oscillations can conspire to produce a spectrum whose areas are close to 2mm, n integral. (Our "pulse area" is m may be any integer, but it is unlikely that the effect could The purpose of the present paper is to point out that under special, but potentially achievable circumstances, the combined temporal behavior of a pulsed laser and an atomic system underassociated with Stark structure, being present only for pulses the power dependence of the effect differs markedly from that (transition probability versus frequency) in multiple photon forfirstion which has a minimum at exact intermediate state

readily be seen for any value but nal. This is discussed in detail below. We designate this splitting as "mock structure" because of its misleading resemblence to Stark structure.

This structure occurs because the probability of inducing the final state is a function of the product of the pulse envelope and the intermediate state population. The latter oscillates in time, and if it should pass through a minimum coincidentally with the maximum in the envelope function of the field strength, the transition probability will be lower than at neighboring frequencies. Where this anticorrelation does not occur.

We consider an isolated atom interacting with a pulsed lawer, undergoing multiple photon ionization from the ground state via a single resonant intermediate state. For simplicity, effects due to the other, nonresonant intermediate states are ignored in this article. The ground and intermediate states are strongly coupled, with the latter connected also to the final states in the continuum. The coupling to the continuum is weak, and is treated perturbatively. With the effect of the continuum on the populations and phases of the intermediate and ground states ignored. Final state amplitudes are to be computed win a quadrature over the intermediate state amplitude.

Let the pulse coupling the two level bound complex have the form f(t) cosin, where f(t) is a smooth function of time, vanishing at t = 2 ... No assumption about the "adiabaticity" or "suddenness"

is made. The time-dependent Schroedinger equation for the amplitudes of this pair of states becomes, in rotating wave approximation,

where & is the detuning of the central frequency of the pulse from exact 15R. The amplitude of a particular continuum state is given perturbatively by

so that the probability of finding the system is state $|k\big>$ as t + + = becomes

5 a 10 that to exp (-; dxt) dt 5 th tesperious of 10 dt. (20)

The total probability of ionization is then obtained by integrating over the continum

(x(t) exp(; 1xt) a= (t) at',

where we have incorporated the density of continum states into the definition of V_{2k} . If the overall process is a two quantum effect, V_{2k} has the same time dependence as V_{12} . If states $|1\rangle$ and $|2\rangle$

differ by one photon, and states $|2\rangle$ and $|x\rangle$ by N-1 photons (for a total absorption of N photons in ionizing the ground state), V_{2k} is an effective operator whose time dependence is $f^{(N-1)}(t)$, Making the usual mild approximations of neglecting the variation of $V_{2k} = V_{0k}^{e,N-1}(t)$ with final state energy and extending the lower limit of the energy integration to — (Geltman 1979), the integrated transition probability becomes

or, since the detuning integral is proportional to 6(t-t'),

$$P = 2\pi (V_0^6)^2 \int_{\mathbb{R}^2} e^{2ir^2} (t) \left| \bullet_2(t) \right|^2 dt.$$
 (5)

We note that while $\mathbf{a}_2(t)$ is a function of the detuning between states $|1\rangle$ and $|2\rangle$, the integral in equation (5) is independent of d_{χ} , the detuning between the intermediate and final states. The population $|\mathbf{a}_2(t)|^2$ undergoes Rabi-type oscillations, whose amplitude in the strong field region is roughly independent of detuning, but where the positions of extrema do depend on d_{χ} the first detuning. Hence, if the time duration of $f^{(23-2)}(t)$ is short, one may expect to see an oscillatory variation of P with laser frequency. This is nost clearly demonstrated in the limiting case of large N, where $f^{(23-2)}$ is approximated by the delta function G(t), G constant,

We see that in this extreme situation, the photofonization spectrum is simply the population variations of level |2 \ transformed into the frequency denain.

We now examine the question of where and whether one may find a sufficiently narrow (27-2 to be possibly observable in actual systems with realistic pulses. We require the coupling between levels |1> and |2> to be sufficiently strong to render the amplitude of the population oscillations at least roughly independent of detuning, without causing the oscillation frequency to be rapid enough to wash out the effect. The details will, of course, depend upon the pulse shape. To simplify the work, we resort to a model in which f(t) is a hyperbolic secant in time. This provides a mooth carelope which might well resemble true pulses. It also emables one to solve the two level part of the problem scalytically (Rosen and Zener 1952, Robiscoe 1978). Equations (1) become

In terms of the parameters a, b, c, 2, given by $\frac{\sqrt{2}}{4}, b = -\frac{\sqrt{5}}{4}, c = \frac{2}{5} + \frac{105}{57}.$

2 - 2 { tent = 1,

with the initial conditions of - 1 and ag - 0 at t - - -, equation

(5) becomes

$$P = k_{2}(V_{0}^{0})^{2} \int_{0}^{1} \left\{ k_{2}(1-z) \right\}^{3-2} \left[k_{2}(z) \right]^{2} dz, \qquad (6)$$

with a_2 given in terms of the variable I by (Robiscos, Roses and Lenes)

where 71 is a hypergeometric function.

The appropriate parameters to characterize the problem is this regime of small V_0^2 are 7, 3, and V_0^7 , where the pulse area V_0^7 determines verther the transition from level $|1\rangle$ to level $|2\rangle$ is saturated, and, if so, the number of cacillations that the population of the intermediate state undergoes for all times.

To gain a quantitative understanding of bow the frequency spectrum for multiphoton ionitation varies with the parameters of the problem, we integrated equation (8) numerically, varying 8 between 3 and 13, and let V₀T assume values in the vicinity of 2m, with n = 1, 2, 3. (At exact resonance, this condition assumes that the amplitude 4₂ passes through zero at 100.) We also explored the regions where the pulse area was very different from an istegral multiple of 2m. We found the following results. For V₀T< 5, the spectra peaked at 500 and distinished monotonically with increasing detuning for all values of 8 in the range investigated, i.e., no structure is predicted. For V₀T = 2m, and 8 ≤ k, no structure

of course, at 2v. For V, T > T, the effect washes out. It reappears was perceived, but for 50%, we found that the calculated forization perceivable in the range from about VoT a 5.5 to VoT a T. peaking, greated nove regidly with increasing I. The excess probability R for high order processes, as one might expect, the effect is nore still greater detuning. As I increased, the value of & for which pail but finite detuning, and then decreased monotonically with Table I shows, for Yor - 3r, the position of the maximum and the the maximus occurred tended to increase slowly. The ratio, 102, warfed from a few percent at Jes to nearly unity at Jell, 1.e., pronounced. It was not necessary that the pulse area be exactly ratio 1-8 of the maximum ionization probability to the resonant probability had a local minimum at 4 . 0, rose to a naximum for in the limit of large I for pulse areas near br. (See belov.) of the maximum fonfamilion probability to the value at 0=0, in-In the the moot structure. We found that the splitting was probability, for fact saing I.

For Fell, we found that some small structure (R = a few percent) could be seen in the wicinity of V_0 = k_T . It appears that this "second harmonic" will be obtainable experimentally only with great difficulty.

While our quantitative predictions are strictly valid only for the hyperbolic secant palse, we believe that qualitatively similar results should obtain for general pulse shapes and be observable with mode-locked lasers.

resonant (MR) intermediate states wis an effective operator approach. which they solved the two level part of the problem in closed form, portion. Their results for smoothly varying pulses do not predict ionization has previously been considered by Grance and Feneralize Interference between the M amplitude and that of the LM has the Book structure, apparently because the N values chosen were below The influence of pulse areas on the spectrum of sultiphoton effect of chifting the maximum of the spectrum, and removing its secunts), for which they used numerical methods in the two level not eliminate, any mock structure effects that might be present. symmetry with respect to detuning. It should also distort, but occur, but also several different smooth pulses (not hyperbolic Grande and Peneutille analyzed not only rectangular pulses, for (1977s, 1977b). These suthors also include the effect of nonand where the doubling predicted in the present article cannot the required minima, according to the present calculation.

That the effect is appreciable only for pulses whose area is near 20 suggests that it may be useful in neutraling the integrated intensity of focused short pulses, provided the resonant dipole matrix element is accurately known. The presence of this doubler would then establish the product V₀T to within rather marrow limits. One could determine whether the effect was indeed most structure by observing if the splitting disappeared with increased and decreased intensity.

To summarize, we have formulated a theory of resonant sultilphoton fontaxtion by pulses valch predicts a spectrom whose maximum
is split into a doublet by purely temporal factors. It appears
similar to the AG Stark effect, but occurs under conditions where
the latter is absent, and exhibits a totally different intensity
dependence.

"Nork supported by the U.S. Office of Maral Research

Position and emplitude of mock structure marine

1	1.02	7.10	1.8	1.31	1.43	1.55	1.66	1.79	1.91	
Zec (19)	2.0	3.0	3.5	6.0	6.4	£.3	57	\$.5	* .5	
.7	•	9		•	6	93	я	a	a	

8 * number of photons needed to louize the ground state. (MT) DAX * position of mock structure maximum in dimensionless units. T is the characteristic time that appears in the hyperbolic secant enrelope function. LtR * ratio of transition probability at maximum to probability at exact resonance.

-75-

References - (continued)

Eogan R M. For R A Robinson E J and Burnham G T 1972 Bull. As. Prive.

Eog. 16 1411

Luger A D and Robinson Z J 1970 Phys. Lett. 31A 247

Lurain M and Mainfray G 1972 Phys. Lett. 32A 1

Robinson Z J and Geltman S 1967 Phys. Rev. 153 4

Robinson Z J and Geltman S 1967 Phys. Rev. 153 4

Robinson Z J and Geltman S 1967 Phys. Rev. 153 4

Robinson Z J and Isenor N R 1968 Can. J. Phys. 46 1537

Zernix W 1964 Phys. Rev. 135 A51

Zernix W 1964 Phys. Rev. 135 A51

Zernix W and Klopfenstein R W 1965 J. Math. Phys. 6 262

Perences

Eall J L. Robinson E J and Branscomb L M 1965 Phys. Rev. Lett. 14 1013 Grannezan E E A and van der Wiel M J 1975 J. Phys. B: Aton. Molec. Fox R A, Kogan R M and Robinson E J 1971 Phys. Rev. Lett. 26 1416 Flank Y, Laplanche G, Jaouen M and Rachman A 1976 J. Phys. B; Agostini P. Lurain M, and Mainfray 6 1971 Pays. Lett. 354 21 Crance M and Feneuille S 1977a J. Phys. (Paris) 27 L-333 Beers B L and Armstrong, Jr. L 1975 Phys. Rev. A12 2447 Chin S L, Isenor N R and Young M 1969 Phys. Rev. 188 7 Dixit S N and Lambropoulos P 1979 Phys. Rev. A19 1576 Eberly J H and O'Well S V 1979 Phys. Rev. A19 1161 Contier Y and Trabin M 1979 Phys. Rev. A 19 264 Chin S L and Isenor N R 1967 Phys. Rev. 158 93 Plank Y and Rechman A 1975 Phys. Lett. 52A 247 Eall J L 1966 IEEE J. Quant. Electr. CT-2 361 Bebb H B and Gold A 1966 Phys. Rev. 143 -1979 Bull. Am. Phys. Soc. 24 66 Geltman S 1963 Phys. Lett. 4 168 Bebb H B 1966 Phys. Rev. 119 25 -1977 Privs. Rev. A 16 1587 -1965 Prys. Lett. 19 616 --- 1967 Phys. Rev. 153 23 Atom Molec. Phys. 9 1409

- 13 -

